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NEWS 4 OCT 07 Multiple databases enhanced for more flexible patent
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NEWS 5 OCT 22 Current-awareness alert (SDI) setup and editing
enhanced
NEWS 6 OCT 22 WPIDS, WPINDEX, and WPIX enhanced with Canadian PCT
Applications
NEWS 7 OCT 24 CHEMLIST enhanced with intermediate list of
pre-registered REACH substances
NEWS 8 NOV 21 CAS patent coverage to include exemplified prophetic
substances identified in English-, French-, German-,
and Japanese-language basic patents from 2004-present
NEWS 9 NOV 26 MARPAT enhanced with FSORT command
NEWS 10 NOV 26 MEDLINE year-end processing temporarily halts
availability of new fully-indexed citations
NEWS 11 NOV 26 CHEMSAFE now available on STN Easy
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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 18:28:28 ON 10 DEC 2008

=> file reg

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

0.21

0.21

FILE 'REGISTRY' ENTERED AT 18:28:39 ON 10 DEC 2008
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STRUCTURE FILE UPDATES: 9 DEC 2008 HIGHEST RN 1082653-47-1
DICTIONARY FILE UPDATES: 9 DEC 2008 HIGHEST RN 1082653-47-1

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TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.

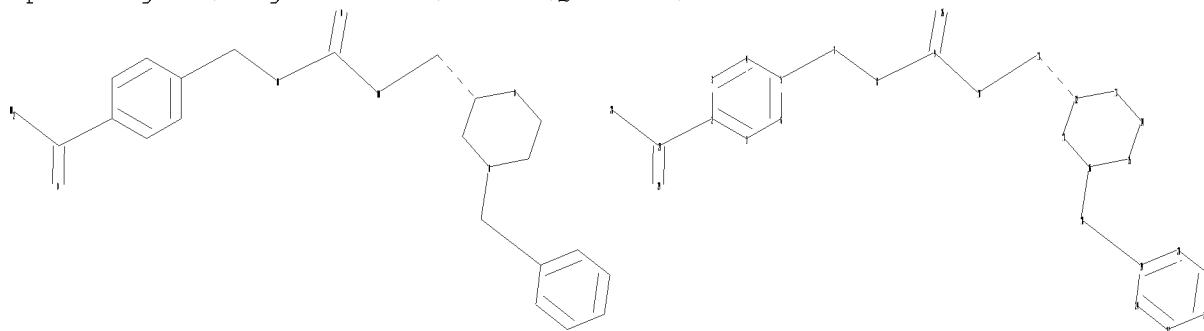
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<http://www.cas.org/support/stngen/stndoc/properties.html>

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Uploading C:\Program Files\STNEXP\Queries\10509521a.str



chain nodes :
7 8 9 10 11 18 26 27 28 29
ring nodes :
1 2 3 4 5 6 12 13 14 15 16 17 19 20 21 22 23 24
chain bonds :
2-27 5-7 7-8 8-9 9-10 9-26 10-11 11-12 16-18 18-19 27-28 27-29
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 12-13 12-17 13-14 14-15 15-16 16-17 19-20
19-24 20-21 21-22 22-23 23-24

exact/norm bonds :
7-8 8-9 9-10 9-26 10-11 11-12 12-13 12-17 13-14 14-15 15-16 16-17 16-18
27-28 27-29
exact bonds :
2-27 5-7 18-19
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6 19-20 19-24 20-21 21-22 22-23 23-24
isolated ring systems :
containing 1 : 12 : 19 :

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:CLASS
11:CLASS 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:CLASS 19:Atom
20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 26:CLASS 27:CLASS 28:CLASS 29:CLASS

L1 STRUCTURE UPLOADED

=> s ll sss full
FULL SEARCH INITIATED 18:29:05 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 306 TO ITERATE

100.0% PROCESSED 306 ITERATIONS 8 ANSWERS
SEARCH TIME: 00.00.01

L2 8 SEA SSS FUL L1

=> file capl
COST IN U.S. DOLLARS SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST 178.36 178.57

FILE 'CAPLUS' ENTERED AT 18:29:08 ON 10 DEC 2008
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FILE COVERS 1907 - 10 Dec 2008 VOL 149 ISS 24
FILE LAST UPDATED: 9 Dec 2008 (20081209/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>

=> s 12

L3 5 L2

=> d 13 1-5 ibib hitstr

L3 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:668939 CAPLUS

DOCUMENT NUMBER: 149:1460

TITLE: Histamine H1 receptor antagonists and CCR3 antagonists for prophylaxis and therapy of allergic diseases

INVENTOR(S): Suda, Makoto; Nanki, Aya

PATENT ASSIGNEE(S): Kowa Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 8pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2008127365	A	20080605	JP 2006-316495	20061124
PRIORITY APPLN. INFO.:			JP 2006-316495	20061124

IT 408303-43-5

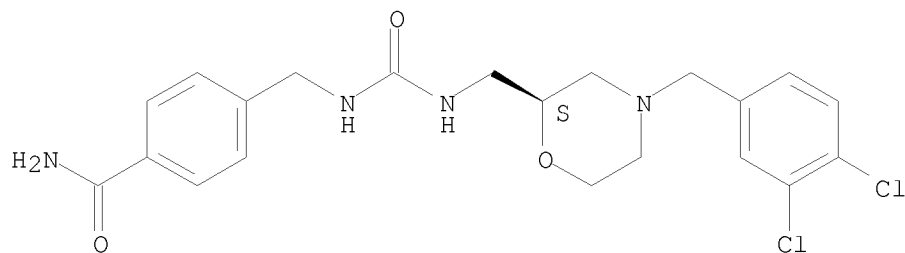
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(histamine H1 receptor antagonists and CCR3 antagonists for prophylaxis and therapy of allergic diseases)

RN 408303-43-5 CAPLUS

CN Benzamide, 4-[[[[(2S)-4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl)methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.



L3 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:796674 CAPLUS

DOCUMENT NUMBER: 139:307774

TITLE: Process for the preparation of aminoalkylmorpholines from hydroxyalkylamines and aminoalkylepoxides.

INVENTOR(S): Hayes, Martin Alistair; Mills, Gail; Swanson, Stephen; Walker, Andrew John; Wilkinson, Mark

PATENT ASSIGNEE(S): Glaxo Group Limited, UK

SOURCE: PCT Int. Appl., 35 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003082835	A1	20031009	WO 2003-EP3343	20030327
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2479819	A1	20031009	CA 2003-2479819	20030327
AU 2003226760	A1	20031013	AU 2003-226760	20030327
EP 1487809	A1	20041222	EP 2003-745297	20030327
EP 1487809	B1	20080109		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003008720	A	20050104	BR 2003-8720	20030327
CN 1642928	A	20050720	CN 2003-807386	20030327
JP 2005532281	T	20051027	JP 2003-580303	20030327
AT 383349	T	20080115	AT 2003-745297	20030327
ES 2298539	T3	20080516	ES 2003-745297	20030327
IN 2004KN01175	A	20060512	IN 2004-KN1175	20040813
ZA 2004006548	A	20050919	ZA 2004-6548	20040817
NO 2004004036	A	20040929	NO 2004-4036	20040924
MX 2004PA09457	A	20050125	MX 2004-PA9457	20040928
US 20050222147	A1	20051006	US 2005-509519	20050502
PRIORITY APPLN. INFO.:			GB 2002-7450	A 20020328
			GB 2000-23902	A 20000929
			GB 2001-7644	A 20010327
			WO 2003-EP3343	W 20030327

OTHER SOURCE(S): MARPAT 139:307774

IT 610785-36-9P

RL: SPN (Synthetic preparation); PREP (Preparation)
(process for the preparation of aminoalkylmorpholines from hydroxyalkylamines and aminoalkylepoxides)

RN 610785-36-9 CAPLUS

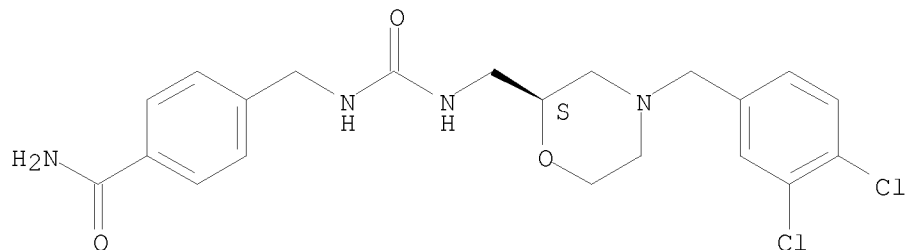
CN Benzamide, 4-[[[[(2S)-4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]-, benzenesulfonate (1:1)
(CA INDEX NAME)

CM 1

CRN 408303-43-5

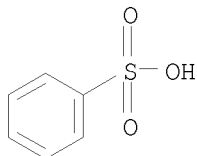
CMF C21 H24 Cl2 N4 O3

Absolute stereochemistry.



CM 2

CRN 98-11-3
CMF C6 H6 O3 S



REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:796495 CAPLUS

DOCUMENT NUMBER: 139:307771

TITLE: Preparation of
4-[[[[(2S)-4-(3,4-dichlorobenzyl)morpholin-2-yl]methyl]amino]carbonyl]amino]methyl]benzamide benzenesulfonate dihydrate and related solvates as a CCR-3 antagonists for the treatment of inflammatory conditions

INVENTOR(S): Cook, John Spencer; Landon, Robert Philip; Walker, Andrew John; Wilkinson, Mark

PATENT ASSIGNEE(S): Glaxo Group Limited, UK

SOURCE: PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003082293	A1	20031009	WO 2003-EP3345	20030327
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2479912	A1	20031009	CA 2003-2479912	20030327
AU 2003226761	A1	20031013	AU 2003-226761	20030327
EP 1487456	A1	20041222	EP 2003-745298	20030327
EP 1487456	B1	20060215		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003008480	A	20050118	BR 2003-8480	20030327
CN 1642554	A	20050720	CN 2003-807364	20030327
JP 2005526808	T	20050908	JP 2003-579830	20030327
AT 317698	T	20060315	AT 2003-745298	20030327
ES 2258724	T3	20060901	ES 2003-745298	20030327
IN 2004KN01286	A	20051230	IN 2004-KN1286	20040902
ZA 2004007463	A	20051012	ZA 2004-7463	20040916

MX 2004PA09456	A	20050125	MX 2004-PA9456	20040928
NO 2004004323	A	20041027	NO 2004-4323	20041012
US 20060089497	A1	20060427	US 2005-509521	20050610
PRIORITY APPLN. INFO.:			GB 2002-7432	A 20020328
			WO 2003-EP3345	W 20030327

OTHER SOURCE(S): MARPAT 139:307771

IT 610785-36-9P

RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of dichlorobenzylmorpholinylmethylaminocarbonylaminomethylbenzamide benzenesulfonate dihydrate and related solvates as CCR-3 antagonists for the treatment of inflammatory conditions)

RN 610785-36-9 CAPLUS

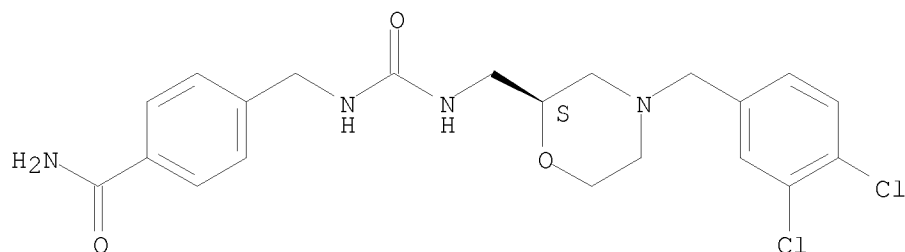
CN Benzamide, 4-[[[[(2S)-4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]-, benzenesulfonate (1:1) (CA INDEX NAME)

CM 1

CRN 408303-43-5

CMF C21 H24 Cl2 N4 O3

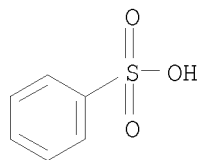
Absolute stereochemistry.



CM 2

CRN 98-11-3

CMF C6 H6 O3 S



IT 408303-43-5P

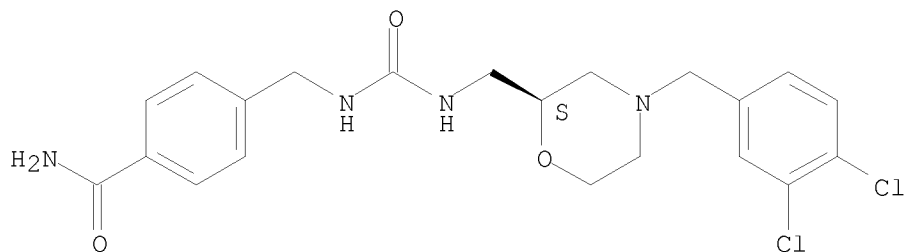
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of dichlorobenzylmorpholinylmethylaminocarbonylaminomethylbenzamide benzenesulfonate dihydrate and related solvates as CCR-3 antagonists for the treatment of inflammatory conditions)

RN 408303-43-5 CAPLUS

CN Benzamide, 4-[[[[(2S)-4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:796494 CAPLUS

DOCUMENT NUMBER: 139:307770

TITLE: Preparation of aralkylureidomorpholines as CCR-3 antagonists for the treatment of inflammatory conditions

INVENTOR(S): Ancliff, Rachael Ann; Cook, Caroline Mary; Eldred, Colin David; Gore, Paul Martin; Harrison, Lee Andrew; Hayes, Martin Alistair; Hodgson, Simon Teanby; Judd, Duncan Bruce; Keeling, Suzanne Elaine; Lewell, Xiao Qing; Mills, Gail; Robertson, Graeme Michael; Swanson, Stephen; Walker, Andrew John; Wilkinson, Mark

PATENT ASSIGNEE(S): Glaxo Group Limited, UK

SOURCE: PCT Int. Appl., 61 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003082292	A1	20031009	WO 2003-EP3340	20030327
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
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EP 1487455	A1	20041222	EP 2003-745296	20030327
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
JP 2005526807	T	20050908	JP 2003-579829	20030327
PRIORITY APPLN. INFO.:			GB 2002-7436	A 20020328
			WO 2003-EP3340	W 20030327

OTHER SOURCE(S): MARPAT 139:307770

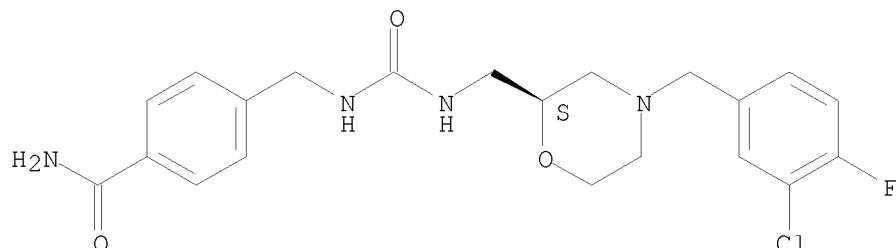
IT 610799-33-2P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of aralkylureidomorpholines as CCR-3 antagonists for the treatment of inflammatory conditions)

RN 610799-33-2 CAPLUS
 CN Benzamide, 4-[[[[(2S)-4-[(3-chloro-4-fluorophenyl)methyl]-2-morpholinyl)methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2002:256246 CAPLUS

DOCUMENT NUMBER: 136:294836

TITLE: Preparation of morpholinylmethylureas for the treatment of inflammatory diseases.

INVENTOR(S): Ancliff, Rachael Anne; Cook, Caroline Mary; Eldred, Colin David; Gore, Paul Martin; Harrison, Lee Andrew; Hodgson, Simon Teanby; Judd, Duncan Bruce; Keeling, Suzanne Elaine; Lewell, Xiao Qing; Robertson, Graeme Michael; Swanson, Stephen

PATENT ASSIGNEE(S): Glaxo Group Limited, UK

SOURCE: PCT Int. Appl., 86 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002026723	A1	20020404	WO 2001-GB4350	20010928
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CA 2423305	A1	20020404	CA 2001-2423305	20010928
AU 2001090146	A	20020408	AU 2001-90146	20010928
BR 2001014321	A	20030701	BR 2001-14321	20010928
EP 1324991	A1	20030709	EP 2001-970027	20010928
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R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
HU 2003003107	A2	20040301	HU 2003-3107	20010928
JP 2004509953	T	20040402	JP 2002-531107	20010928
NZ 525055	A	20040924	NZ 2001-525055	20010928
EP 1586567	A1	20051019	EP 2005-76503	20010928
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IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

AT 345331	T	20061215	AT 2001-970027	20010928
ES 2275729	T3	20070616	ES 2001-970027	20010928
ZA 2003002461	A	20040628	ZA 2003-2461	20030327
NO 2003001443	A	20030526	NO 2003-1443	20030328
MX 2003PA02813	A	20031015	MX 2003-PA2813	20030328
IN 2003KN00419	A	20050311	IN 2003-KN419	20030408
US 20040058907	A1	20040325	US 2003-381871	20030721
US 7157457	B2	20070102		

PRIORITY APPLN. INFO.:

GB 2000-23973	A	20000929
GB 2001-7643	A	20010327
EP 2001-970027	A3	20010928
WO 2001-GB4350	W	20010928

OTHER SOURCE(S): MARPAT 136:294836

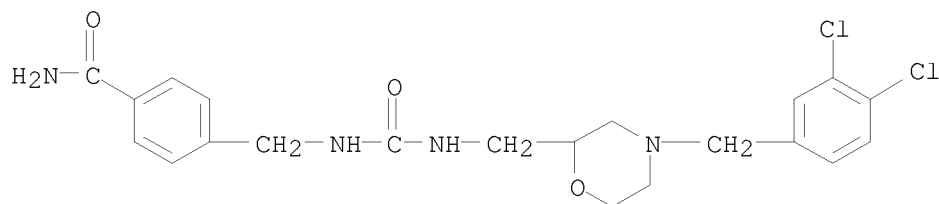
IT 408303-42-4P 408303-43-5P 408303-44-6P
 408304-13-2P 408304-74-5P 408304-75-6P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of morpholinylmethylureas for the treatment of inflammatory diseases)

RN 408303-42-4 CAPLUS

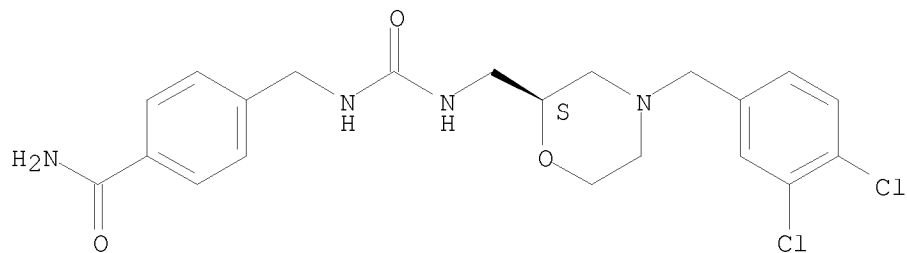
CN Benzamide, 4-[[[[[4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)



RN 408303-43-5 CAPLUS

CN Benzamide, 4-[[[[[[(2S)-4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)

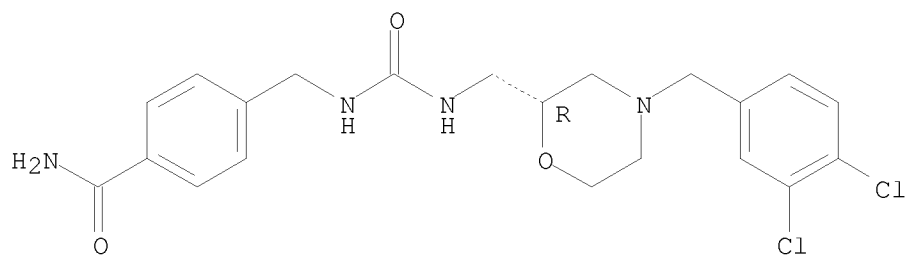
Absolute stereochemistry.



RN 408303-44-6 CAPLUS

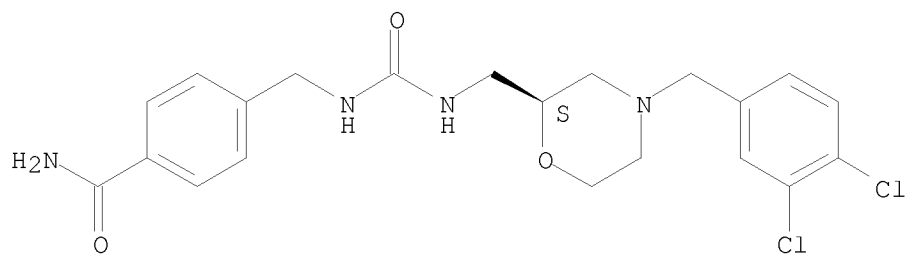
CN Benzamide, 4-[[[[[[(2R)-4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.



RN 408304-13-2 CAPLUS
 CN Benzamide, 4-[[[[(2S)-4-[(3,4-dichlorophenyl)methyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]-, hydrochloride (1:1) (CA INDEX NAME)

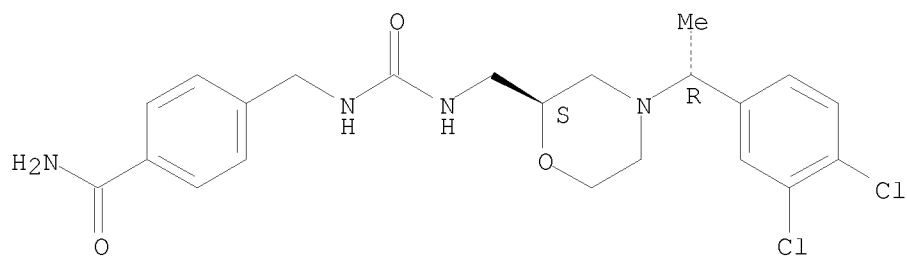
Absolute stereochemistry.



● HCl

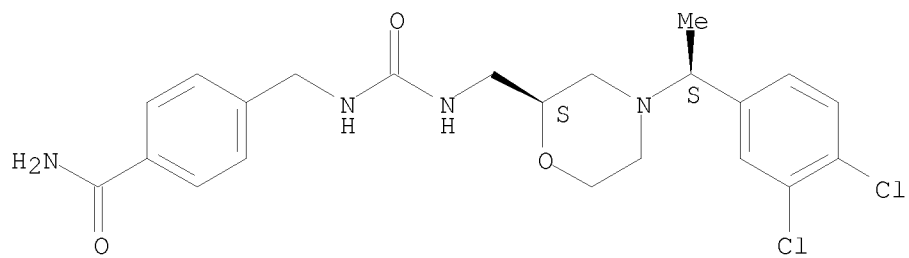
RN 408304-74-5 CAPLUS
 CN Benzamide, 4-[[[[(2S)-4-[(1R)-1-(3,4-dichlorophenyl)ethyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.



RN 408304-75-6 CAPLUS
 CN Benzamide, 4-[[[[(2S)-4-[(1S)-1-(3,4-dichlorophenyl)ethyl]-2-morpholinyl]methyl]amino]carbonyl]amino]methyl]- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT:

10

THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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